## **AMENDMENTS TO THE CLAIMS:**

This listing of claims will replace all prior versions and listings of claims in the application:

1. (Original) A cyclic carbonate-containing polymeric compound represented by formula (I):

$$\begin{array}{c|c}
CH_{2} & CH_{2} & CH_{2} \\
CH_{2} & CH_{2} & CH_{2} \\
CH_{2} & CH_{2} & CH_{2}
\end{array}$$

$$\begin{array}{c|c}
CH_{3} & CH_{2} \\
CH_{2} & CH_{2}
\end{array}$$

$$\begin{array}{c|c}
CH_{3} & CH_{2} \\
CH_{2} & CH_{2}
\end{array}$$

$$\begin{array}{c|c}
CH_{3} & CH_{2}
\end{array}$$

$$\begin{array}{c|c}
CH_{3} & CH_{2}
\end{array}$$

$$\begin{array}{c|c}
CH_{3} & CH_{2}
\end{array}$$

wherein p, q, and r independently represent the molar composition ratio of each monomer unit: p is a number over 0; q and r are each a number not smaller than 0; and the sum of p, q, and r is 1 or smaller.

- 2. (Currently Amended) A method for producing the cyclic carbonate-containing polymeric compound according to claim 1 comprising a first step of epoxidizing deproteinized natural rubber and a second step of allowing the epoxidized natural rubber obtained via the first step to react with supercritical carbon dioxide.
- 3. (Original) The method according to claim 2, wherein the second step is carried out in the presence of a polar organic solvent and/or an ionic liquid.

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- 4. (Original) The method according to claim 3, wherein the polar organic solvent is at least one member selected from the group consisting of N,N-dimethylformamide, N,N-diethylformamide, N,N-diethylacetamide, N,N-diethylacetamide, and N-methylpyrrolidone.
- 5. (Original) The method according to claim 3, wherein the ionic liquid is at least one member selected from the group consisting of 3-methyl-1-octylimidazolium tetrafluoroborate, 1-hexyl-3-methylimidazolium tetrafluoroborate, 1-butyl-3-methylimidazolium tetrafluoroborate, 1-ethyl-3-methylimidazolium tetrafluoroborate, 1-ethyl-3-methylimidazolium tetrafluoroborate, 1-tifluoromethanesulfate.
- 6. (Original) The method according to claim 2, wherein the second step is carried out at a reaction temperature between 50° C. and 200° C.
- 7. (Original) The method according to claim 2, wherein the second step is carried out at a supercritical carbon dioxide pressure of between 5 MPa and 20 MPa.
- 8. (Original) The method according to claim 2, wherein the second step is carried out for 0.5 hour to 20 hours.